

This Page Is Inserted by IFW Operations  
and is not a part of the Official Record

## **BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning documents *will not* correct images,  
please do not report the images to the  
Image Problem Mailbox.**



FC

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 :  B08B 7/00, C11D 7/50		A1	(11) International Publication Number: WO 92/09378  (43) International Publication Date: 11 June 1992 (11.06.92)
(21) International Application Number:	PCT/US91/08937	(72) Inventors; and	
(22) International Filing Date:	29 November 1991 (29.11.91)	(75) Inventors/Applicants ( <i>for US only</i> ) :	BOLDEN, Paul, L. [US/US]; 18 Florence Point Drive, Fernandina Beach, FL 32034 (US). HAYES, Michael, E. [US/US]; 71 Oak Lane, Fernandina Beach, FL 32034 (US). BRADDY, John, E. [US/US]; 1567 Penbrook Drive, Fernandina Beach, FL 32034 (US).
(30) Priority data:	620,500 29 November 1990 (29.11.90) US	(74) Agents:	BARRON, Alexis et al.; Synnestvedt & Lechner, 2600 One Reading Center, 1101 Market Street, Philadelphia, PA 19107 (US).
(60) Parent Application or Grant		(81) Designated States:	AT (European patent), BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, KR, LU (European patent), NL (European patent), SE (European patent), US.
(63) Related by Continuation			
US	620,500 (CIP)		
Filed on	29 November 1990 (29.11.90)		
(71) Applicant ( <i>for all designated States except US</i> ): PETRO-FERM INC. [US/US]; 5400 First Coast Highway, Fernandina Beach, FL 32034 (US).		Published	<i>With international search report.</i>

(54) Title: A PROCESS AND COMPOSITION FOR CLEANING CONTAMINANTS WITH TERPENE AND MONOBASIC ESTER

(57) Abstract

An industrial cleaning process in which a contaminant comprising soldering flux and/or adhesive tape residue is removed from a substrate contaminated therewith, for example, a printed circuit board contaminated with rosin flux, comprising: (A) contacting the substrate with a terpene-based cleaning composition which is substantially free of water-soluble organic material, for example, a surfactant, for a period of time sufficient to solubilize the contaminant; (B) removing the composition and solubilized contaminant from the substrate by contact with water having a temperature of about 70 °F to about 140 °F (but preferably lower) for a period of time of no longer than about 10 minutes to provide a substrate having a contamination rating of no greater than about 14 micrograms NaCl equivalent/square inch (MIL-P-28809A), and to thereby form a mixture comprising the composition, contaminant and water; and (C) separating the water from said mixture, the separated water being substantially free of water-soluble organic material and having a chemical oxygen demand of no greater than about 1,000 ppm, and also a cleaning process utilizing a cleaning composition comprising a major amount of a monobasic ester to clean surfaces that are contaminated with an organic residue.

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	ES	Spain	MG	Madagascar
AU	Australia	FI	Finland	ML	Mali
BB	Barbados	FR	France	MN	Mongolia
BE	Belgium	GA	Gabon	MR	Mauritania
BF	Burkina Faso	GB	United Kingdom	MW	Malawi
BG	Bulgaria	GN	Guinea	NL	Netherlands
BJ	Benin	GR	Greece	NO	Norway
BR	Brazil	HU	Hungary	PL	Poland
CA	Canada	IT	Italy	RO	Romania
CF	Central African Republic	JP	Japan	SD	Sudan
CG	Congo	KP	Democratic People's Republic of Korea	SR	Sweden
CH	Switzerland	KR	Republic of Korea	SN	Senegal
CI	Côte d'Ivoire	LJ	Liechtenstein	SU+	Soviet Union
CM	Cameroon	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TG	Togo
DE*	Germany	MC	Monaco	US	United States of America

+ Any designation of "SU" has effect in the Russian Federation. It is not yet known whether any such designation has effect in other States of the former Soviet Union.

A PROCESS AND COMPOSITION FOR CLEANING CONTAMINANTS WITH  
TERPENE AND MONOBASIC ESTER

---

Field of the Invention

This invention relates to the removal of a residue from a surface that is contaminated therewith, for example, to the 5 cleaning of deposits of soldering flux and/or other residues from a substrate which is contaminated therewith. More specifically, this invention relates to the use of a cleaning composition containing an ester. This invention relates also to the use of terpenes, preferably in admixture with an ester, 10 in an improved cleaning process for removing contaminants such as soldering flux and/or adhesive tape residue from a substrate.

The invention will be described initially in connection with its applicability to the cleaning of printed circuit 15 boards. However, the invention has wider applicability, as described hereinbelow.

In the fabrication of printed circuit boards, it is known to use soldering fluxes in order to ensure the firm, uniform bonding thereto of solder which functions to adhere to the 20 boards components thereof. Such flux must be removed from the finished board. If not removed, deposits of the flux, in even trace or minute amounts, can cause the circuitry of the board to fail. An example of a flux that is widely used in the

fabrication of printed circuit boards is rosin flux, a material which is water insoluble.

Various types of materials have been proposed for cleaning deposits of rosin flux from the surface of a printed circuit board. Traditionally, chlorofluorocarbons (CFCs) and other halogenated solvents, including chlorinated hydrocarbon solvents, were used widely. However, these materials are not presently favored because of environmental problems that are associated therewith. Examples of other cleaning materials for removing deposits of rosin flux from substrates contaminated therewith include: (a) alkaline cleaning compounds such as alkanol amines; (b) a mixture of N-methyl-2-pyrrolidine and an alkanol amine (U.S. Patent No. 4,276,186); (c) terpene compounds; and (d) dibasic acid esters (U.S. Patent No. 4,934,391).

Although there are advantages and disadvantages associated with the use of each of the aforementioned type cleaning materials, the use of terpene compounds in cleaning printed circuit boards of flux deposits has become more and more popular because they possess a combination of unusually desirable properties. The present invention is directed to an improved process for cleaning printed circuit boards and other substrates of deposits of soldering flux and/or other residues with a terpene-based cleaning composition which optionally, but preferably, includes an ester.

The present invention is directed also to the use of a non-toxic, environmentally safe cleaning composition comprising an ester and optionally other materials, for example, a terpene.

Reported Developments

In the cleaning operation, deposits of soldering flux are dissolved by the water-insoluble, liquid cleaner, for example, terpene compounds as they are contacted therewith. A review of 5 the related art reveals that special consideration is given to removing from the terpene-treated substrate the dissolved contaminants that adhere thereto for those applications which require that the substrate be highly cleaned. Thus, the effective removal of the dissolved soldering flux and the 10 terpene solvent from the substrate in which the flux is dissolved is an important aspect of the overall cleaning process.

The basic patent on the use of terpenes to clean printed circuit boards of soldering flux is U.S. Patent No. 4,640,719, 15 assigned to the same assignee as the present invention. With respect to the removal from the terpene-treated substrate of dissolved flux and other contaminants, this patent teaches as follows:

"Because they [terpene compounds] are almost completely insoluble in water, however, the terpenes cannot be directly flushed away by water. Alternatively and preferably, the terpene compounds are combined with one or more terpene emulsifying surfactants. The addition of such surfactants facilitates removal of the terpenes from printed wiring boards by rinsing with water, whereby the terpenes are formed into oil-in-water emulsions." 25 (Column 3, lines 54-62)

The '719 patent discloses further that the terpene-based 30 composition can include up to 40 wt.% of the emulsifying surfactant, with a preferred composition containing 10 wt.% of surfactant. Such surfactants are typically water-soluble organic materials.

European patent application bearing publication number 0 35 354 027 A2 discloses cleaning compositions comprising a major amount of selected cyclic terpenes, including para-

menthadienes, and a minor amount of terpene alcohol for use in cleaning a variety of substrates, including circuit boards.

This patent discloses that the compositions can include surfactants and specifically states the following:

5 "It has been found helpful to add surfactants to solvent compositions which are to be used in combination with a water rinse. It has been found that the addition of the surfactant to the cyclic terpene mixture enhances the water rinsability of the 10 solvent composition. Further, it has been found that in dealing with certain types of soil, the addition of a surfactant to the mixture of cyclic terpenes enhances the ability of the cyclic terpene mixture to remove the soil from the substrate."

15 (Column 4, line 57 to Column 5, line 1)

This publication discloses further that the surfactants can be used in concentrations of from 0.1 to 10 wt.%.

Another patent which relates to the removal of soldering flux from substrates such as printed circuit boards is U.S.

20 Patent No. 4,867,800. The cleaning composition which is the subject of this patent comprises a terpene compound and a dibasic ester which has at least 2 wt.% solubility in water at 25°C. This patent discloses that such compositions can be used to achieve improved cleaning relative to the use of terpene-based compositions that include emulsifying surfactants, which the patentee refers to as an unnecessary, but optional ingredient of his composition. The patent discloses also that the terpene-treated surface is preferably rinsed with water, such as deionized water. In an exemplary embodiment disclosed 25 in the patent, the rinse water which was used in the cleaning process had a temperature of about 150°F.

30

A common characteristic of each of the processes described in the aforementioned publications is the required or preferred use of water-soluble organic materials in combination with the 35 terpene constituent of the cleaning composition. While considered essential to achieve high grade cleaning, the presence of such water-soluble organic materials leads to

problems which are associated with other aspects of the overall cleaning process.

One significant problem crops up in connection with disposal of the rinse water that is generated during the 5 cleaning process. The amount of such rinse water can be substantial, with some industrial applications using, for example, two to five gallons of rinse water per minute of operating time. For various industrial applications, it is considered ideal to simply discharge the rinse water into the 10 local sewage system. However, as a result of the use in the cleaning composition of water-soluble materials, such discharge typically runs afoul of environmental standards which require that the chemical oxygen demand (COD) of the discharged water be no greater than some defined value. For example, some 15 communities specify that the COD should be no greater than about 1,000 ppm for industrial water that is discharged into the local sewage system. There are other communities with even higher standards, for example, CODs of no greater than about 300 ppm. However, in using cleaning compositions of the type 20 described above, the presence of the water-soluble materials can result in the rinse water having as much as 10,000 ppm or more of COD. This indeed presents a problem for the fabricators of circuit boards inasmuch as it requires the fabricators to adopt additional steps to treat the water in a 25 manner such that environmental standards are met. This, of course, increases the cost of the overall manufacturing process.

There are reports (unconfirmed) of the use of surfactant-free, terpene-based compositions in applications for removing 30 solder fluxes from various types of substrates. However, it appears that, in such applications, the terpene-treated substrate is rinsed with surfactant-containing water or requires that the substrate be rinsed with either boiling water or near boiling water. This, of course, is undesirable because

of the increased costs associated with the provision of energy that is needed to heat the significant amounts of water that are required for the rinsing operation.

In view of exemplary shortcomings of the present state of  
5 the art related to the use of terpene-based compositions to  
clean soldering flux from the surfaces of printed circuit  
boards, as described above, the present invention is directed  
to an improved, more efficient and economical process which  
avoids or significantly alleviates such shortcomings.

10 As mentioned briefly above, CFCs and other halogenated  
solvents which have been used widely as cleaning agents for  
many years are no longer materials of choice because of their  
detrimental effects on the earth's stratospheric ozone layer.  
Species of these materials are perceived also to be  
15 carcinogenic. Due to their environmental and health threats,  
they are being phased out of production even though they have a  
variety of properties which otherwise make them ideal cleaning  
agents. They are moderately good oil and grease solvents, they  
have no flash points, they evaporate quickly due to their high  
20 vapor pressures, and some have low toxicity.

Alkaline cleaning compounds known as the alkanol amines,  
usually in the form of monoethanolamine, have been used for  
rosin flux removal as an alternative to the toxic halogenated  
hydrocarbon solvents. These compounds chemically react with  
25 rosin flux to form a rosin soap through the process of  
saponification. Other organic substances such as surfactants  
or alcohol derivatives may be added to these alkaline cleaning  
compounds to facilitate the removal of such rosin soap. The  
use of N-methyl-2-pryrrolidone in admixture with a water-  
30 miscible alkanolamine is disclosed in U.S. Patent No.  
4,276,186. Unfortunately, these compounds, like the water  
soluble soldering fluxes, have a tendency to cause corrosion of  
the surfaces and interfaces of printed wiring boards if they

are not completely and rapidly removed during the fabrication process.

Another material which has been identified as an alternative to the use of halogenated solvents is the class of compounds known as dibasic esters. The use of these materials is desirable because they have relatively high flash points and are not skin irritants. However, they have the disadvantage of being moderately water soluble. Water solubility is undesirable because dissolved materials are difficult to separate from the water when used in a process involving water rinse (as is usually the case). Thus, it is difficult to dispose of the water rinse generated in the cleaning process. Closed-loop systems are relatively difficult and/or expensive to install when the cleaning compound is substantially water soluble.

Even terpenes, which have been recognized as environmentally acceptable alternatives to halogenated solvents and which offer good performance in a wide range of cleaning applications, have some disadvantages. They have an objectionable odor, low flash point, and a tendency to irritate skin, lungs and nasal passages.

Still another class of compounds in use is mineral spirits or other petroleum distillates. These materials often have low flash points, with high evaporative losses, and produce dermatitis, headaches and nausea in use. Petroleum distillates also require frequent change-outs in use because their cleaning power is greatly diminished with minor loadings of oil and machine residues.

The present invention encompasses also the provision of a cleaning process involving the use of a cleaning agent which has a combination of properties such that the agent is

particularly suitable for safe and efficient use in a wide variety of cleaning processes.

Summary of The Invention

In accordance with the present invention, it has been  
5 found that a composition comprising a major amount of a monobasic ester can be used effectively in a variety of different types of cleaning applications which heretofore involved the use of halocarbons, alkaline detergents, or other cleaning agents as described above, the use of each of the  
10 aforementioned being accompanied by serious disadvantages. The monobasic esters, which can be used to clean a variety of surfaces, for example, metals, plastics, and electronic assemblies, offer the advantage of water insolubility, low toxicity, low skin irritation, high flash point and low odor.

15 Accordingly, there is provided, in accordance with the present invention, a method for removing an organic residue from a surface contaminated therewith comprising:

(A) contacting the surface with a cleaning composition comprising a major amount of a monobasic acid ester;

20 (B) maintaining the cleaning composition in contact with the surface for a period of time at least sufficient to solubilize the residue; and

(C) removing cleaning composition and solubilized residue from the surface.

25 Examples of monobasic esters that can be used in the cleaning operation are methyl, ethyl, and isopropyl esters of medium chain (about C<sub>6</sub> to about C<sub>18</sub>) monobasic acids.

The conditions under which the monobasic ester composition are used can be typical of conditions used for other organic cleaning solvents. They can be used, for example, at a temperature within the range of room temperature to about 150°F or higher and under conditions which involve a contact time of about 1 to about 10 minutes. Conventional means can be used to apply the monobasic ester composition to the soiled substrates, for example, spray, agitated immersion, and soaking.

Use of the monobasic ester composition of the present invention avoids disadvantages associated with the use of prior art compositions, as noted above. Use of a composition within the scope of the present invention offers the advantages of: (a) good oil solvency; (b) good solvency for solder fluxes; (c) excellent ability to remove lapping compounds, chips, grinding residues and other solid materials from metal surfaces after machining operations; (d) the ability to remove the solids even though highly loaded with cutting oils; (e) extremely low toxicity and skin irritation; (f) low odor or pleasing fragrances; (g) high flash points; and (h) water insolubility. Furthermore, the monobasic ester composition does not have a harmful effect on the earth's stratospheric ozone layer and they generally have a relatively low vapor pressure, and are, therefore, less volatile, providing the advantage of lower volatile organic compound (VOC) pollutant emissions than prior art compositions.

In addition, there is provided in accordance with the present invention an industrial cleaning process in which a contaminant comprising soldering flux and/or adhesive tape residue is removed from a substrate contaminated therewith comprising:

(A) contacting the substrate with a terpene-based cleaning composition which is substantially free of water-soluble organic material for a period of time sufficient to solubilize the contaminant;

5 (B) removing the composition and solubilized contaminant from the substrate by contact with water having a temperature of about 70°F to about 140°F for a period of time of no longer than about 10 minutes to provide a substrate having a contamination rating of no greater than about 14  
10 micrograms NaCl equivalent/square inch (MIL-P-28809A), and to thereby form a mixture comprising the composition, contaminant and water; and

15 (C) separating the water from said mixture, the separated water being substantially free of water-soluble organic material and having a chemical oxygen demand of no greater than about 1,000 ppm.

With respect to Step (A) above, the following procedure is used to determine the content of water-soluble organic material in a terpene-based composition. Equal volumes of water and the  
20 terpene-based composition are combined and allowed to separate into two distinct phases or layers at about 70°F. To the extent that the terpene-based composition contains water-soluble organic material, it will migrate to the water layer. The COD of the water layer is then determined according to  
25 standard technique. When the water layer has a COD of no greater than about 1,000 ppm, the terpene-based composition is considered to be "substantially free of water-soluble organic material".

Thus, in accordance with the present invention, it has  
30 been found, contrary to the teachings of the prior art, that it is possible to produce a highly clean surface by the use of a terpene-based cleaning composition which is free of or essentially free of emulsifying surfactants and/or other water soluble organic materials.

And such highly cleaned surfaces can be produced under practical operating conditions, for example, at ambient temperature or relatively low elevated temperature and within times and water contact conditions that are presently used in industrial applications. And of great advantage is the fact that the water collected from the cleaning process has a sufficiently low COD to enable it to be lawfully discharged to a sewage system without purification or other treatment. Alternatively, the collected water can be recycled for use in the cleaning process, as described hereinbelow.

With respect to the high degree of cleaning that can be achieved by use of the present invention, the present cleaning process, including either the monobasic ester-based composition or the terpene-based composition, can be used to produce levels of residual contamination that are equal to or far below the military requirement of 14 micrograms NaCl equivalent/in<sup>2</sup> (MIL-P-28809A) (see the aforementioned '719 patent, column 7, lines 18 to 32 for additional information concerning the evaluation of the degree of cleanliness of substrates based on the involved military test). A specific example of the degree of cleanliness that can be achieved by practice of the present invention is set forth in the Example section hereinbelow.

#### Detailed Description of the Invention

##### The Monobasic Ester Development

The cleaning composition of the present invention comprises a major amount of a monobasic ester compound, that is, the ester compound comprises in excess of 50 wt.% of the composition, based on the total weight of the composition. The ester compound, which is a liquid at room temperature, can be represented by the following formula:



wherein R and R<sub>1</sub> are hydrocarbon groups, which may be the same or different, in which the sum of the carbon atoms in R and R<sub>1</sub> is at least about 8. The hydrocarbon groups can be aliphatic or aromatic, substituted or unsubstituted. Examples of 5 aromatic groups include aryl groups, for example, phenyl and benzyl. In preferred form, each of R and R<sub>1</sub> is an alkyl group with the total number of carbon atoms in R and R<sub>1</sub> being no greater than about 20. The alkyl groups can be straight or branched chain and unsubstituted or substituted, for example, 10 with groups such as hydroxyl and ether groups. The cleaning composition can comprise two or more of the monobasic esters.

Particularly preferred monobasic esters for use in the practice of the present invention are those in which the total number of carbon atoms in R and R<sub>1</sub> is about 9 to about 17, with 15 R being an alkyl group having 1 to about 4 carbon atoms and R<sub>1</sub> being an alkyl group having about 8 to about 16 carbon atoms. Particularly preferred monobasic esters are methyl laurate and isopropyl laurate, with methyl caprate and isopropyl myristate being even more preferred. The preferred esters are excellent 20 solvents for a variety of different kinds of organic residues that typically contaminate the surfaces of a variety of types of materials.

Examples of commercially available monobasic esters which can be used in the practice of the present invention include: 25 methyl caprylate, methyl caprate, methyl laurate, methyl palmitate, isopropyl myristate, isopropyl laurate and isopropyl stearate.

It will be appreciated that a particular carboxylic monobasic ester can be synthesized readily by available 30 methods. For example, monobasic esters can be prepared by reacting an alkyl alcohol and a fatty acid, for example, a C<sub>1</sub> to about a C<sub>8</sub> alkanol and about a C<sub>3</sub> to about a C<sub>18</sub> fatty acid.

The insolubility of the ester is determined by the following procedure which is conducted at about 70°F. Equal volumes of water and the ester are combined and allowed to separate into two distinct phases or layers. The chemical 5 oxygen demand (COD) of the water layer is then determined according to standard technique. If the water layer has a COD of no greater than about 5,000 ppm, the ester is considered to be "substantially insoluble in water". Preferably the 10 solubility characteristics of the ester are such that the water layer has a COD of no greater than about 1,000 ppm - such esters being referred to as "water-insoluble".

Other preferred properties of the ester are that it have a flash-point of greater than about 140°F, preferably greater 15 than about 200°F, and low odor. In addition, the ester should be non-irritating to the skin and have good chemical stability.

The cleaning composition of the present invention can comprise 100 wt.% of the monobasic ester. However, there are applications in which it will be advantageous to include other materials in the composition, but in an amount less than 50 20 wt.%. For example, some types of residues can be removed more readily by using other cleaning agents, including, for example, organic or inorganic solvents, in admixture with the monobasic ester. In preferred form, such other cleaning agents should be miscible with the monobasic ester. Examples of other cleaning 25 agents that can be used include terpenes, for example, para-menthadienes, dibasic esters, petroleum hydrocarbons, ketones, ethers, and alcohols. Preferred cleaning agents for use in combination with the monobasic ester are terpenes, ethers, and alcohols.

30 For applications in which it is desirable to use another cleaning agent in combination with the monobasic ester, the amount thereof can be selected based on the results desired. For most applications, it is believed that the composition will

comprise about 55 to 90 wt.% of the monobasic ester and about 10 to about 45 wt.% of the other cleaning agent.

The organic-based cleaning composition of the present invention can include also a surfactant for promoting 5 the removal of residual cleaning composition and contaminant from the substrate being cleaned. In some applications, the presence of surfactant aids also in the cleaning of contaminant from the substrate.

10 The surfactant can be a nonionic, anionic, cationic or amphoteric surfactant. It is preferred to use a nonionic surfactant, for example, condensation products of alkylene oxides, for example, ethylene oxide, with any of a variety of hydrophobic moieties, as are well known in the art. Examples of these products are ethoxylated aliphatic alcohols, 15 ethoxylated alkylphenols, ethoxylated amines, amides, and the like. Sulfated, sulfonated, phosphated and carboxylated hydrophobes, such as alcohols, can also be used, as can the salts of each. Particularly preferred surfactants are ethoxylated aliphatic alcohols and their derivatives.

20 Examples of surfactants that can be used are ethoxylated nonylphenols having about 4 to about 10 ethylene oxide groups and detergent alcohols, for example, those comprising about C<sub>9</sub> to about C15 linear alcohols and about 3 to about 10 ethylene oxide groups.

25 The proportion of surfactant comprising the composition will vary, depending on the particular ester or combination of esters used and the nature of the particular cleaning application. Generally the surfactant will comprise about 2 to 30 about 20 wt% of the composition. In a highly preferred embodiment, the composition comprises a mixture of about 8 to about 12 wt.% of surfactant and about 88 to about 92 wt.% of monobasic ester.

The ester cleaning composition of this invention can be used in a wide variety of applications because the ester cleaning agent is capable of dissolving a wide variety of organic materials that typically contaminate substrates and it  
5 is compatible (not harmful) with many types of materials that comprise substrates that require cleaning. Thus, the ester-based composition can be used in removing cutting oils, greases, greasy dirt, solder fluxes, waxes, tape residues, coolants, lubricants, lapping compounds, cuttings turnings,  
10 chips and other machining residues from machined parts and for hard surface cleaning in general. They can be used advantageously to clean metals, plastics and electronic assemblies.

The ester cleaning composition of the present invention  
15 can be used in cleaning effectively printed circuit boards, as well as any type of substrate which is contaminated with soldering flux and/or other types of residue that typically contaminate the surfaces of printed circuit boards, for example, adhesive tape residue, waxes and fingerprints. Thus,  
20 printed wiring boards and other articles which require the use of soldering flux in the making of mechanical, electromechanical or electronic connections with metal can be cleaned in accordance with the present invention. In the fabrication of such articles, rosin soldering flux is used  
25 widely. However, there also can be used other types of soldering fluxes, including, for example, synthetic activated fluxes, organic acid-based fluxes and fluxes contained within soldering paste.

With respect to the degree of substrate cleanliness that  
30 can be achieved by the practice of the present invention, levels of contamination that meet or are below the military specification MIL-P-28809A requirement of 14 micrograms NaCl equivalent/square inch can be achieved. The ester cleaning

composition has been used effectively to provide substrates having a contamination level below about 10 micrograms NaCl equivalent/square inch.

The contaminated surface should be contacted with the  
5 cleaning composition for a period of time sufficient to solubilize the contaminant. The time for effecting solubilization will depend on the temperature of the cleaning composition and the means by which it is applied. The temperature of the composition can be like that typically used  
10 for conventional cleaning compositions, for example, room or ambient temperature (about 70°F) to about 150°F. For some applications, for example, wax removal, even higher temperatures can be used, for example, up to about 240°F.  
Exemplary means for applying the cleaning composition are  
15 immersion, brushing, and spraying, for example, spraying in air or beneath the surface of the cleaning composition. It is believed that spraying the cleaning composition onto the substrate will be used most widely. For batch operations, typical contact times are about 1 to about 10 minutes. For  
20 continuous operations, typical contact times are about 30 seconds to about 3 minutes.

The solubilized contaminant and residual cleaning composition can be removed from the substrate by any suitable means, for example, by air blasting, rinsing with water or an  
25 organic rinsing agent, or by wiping, for example, with an absorbent material. It should be understood that, for some applications, it is not necessary to remove completely the cleaning composition and the contaminant from the substrate. In such an application, the cleaning composition is simply  
30 allowed to drain from the surface.

Drying of the substrate is optional. If it is desired to dry the surface, drying can be accelerated by the use of heat. For example, the substrate can be dried in an oven and/or it

can be dried with forced air, for example, air heated to 100°F or above.

In accordance with this invention, it has been found that the monobasic ester cleaning composition can be used effectively to clean a plurality of parts even after prolonged use with build up in the composition of a relatively high amount of dissolved organic contaminants that have been removed from the treated parts. In one application, it was observed that a bath of the cleaning composition continued to clean effectively parts covered with an oily material even though the bath contained about 40 wt.% of the dissolved organic residue. Accordingly, the present invention encompasses also the continued use of the monobasic ester composition to clean organic residues from the surfaces of substrates contacted therewith as dissolved organic residue builds up in concentration in the composition to a relatively high amount, but preferably to an amount no greater than about 30 wt.%, and more preferably to an amount of about 20 wt.% (based on the total weight of the cleaning composition and the amount of dissolved organic contaminant).

#### Examples of the Monobasic Ester Development

Examples which follow are illustrative of the present invention. Comparative examples are set forth also.

The first example is a comparative example which shows the use of CFC-113 (hereinafter "the CFC composition"), an example of a prior art cleaning composition comprising 1,1,2-trichloro-trifluoroethane. This comparative example is followed by an example of the present invention showing the use of a cleaning composition containing a monobasic ester to clean the same type of machine parts as were cleaned by the use of CFC composition.

#### Example C-1

In machining operations during the manufacture of component parts of automotive fuel injectors, the parts being machined are subjected to statistical in-process quality control measurements for dimensional conformity. The parts 5 were taken from the machine and swished by hand in a container of the CFC composition to remove cutting oil, chips and grinding residues. The chips and grinding residues settled to the bottom of the container and the cutting oil dissolved in the CFC composition. The parts were blown off with a jet of 10 air to remove the residual CFC composition. The parts were then placed in a laser gauge to measure dimensional conformity.

Example 1

A cleaning composition comprising 95 wt.% isopropyl myristate and 5 wt.% of other unidentified monobasic esters 15 (by-products of the manufacturing process) hereafter "the ester composition" was substituted for the CFC-113 composition. The fuel injectors were taken from the machine and swished in the container of the ester composition. The machine operators were in direct contact with the ester composition during this 20 cleaning operation. The chips and grinding residues settled to the bottom of the container and the cutting oil dissolved in the ester composition. The parts were blown off with a jet of air to remove the residual ester composition and then placed in the laser gauge to measure dimensional conformity.

25 The measurements in the laser gauge were the same, and performed in the same manner, as those associated with the parts cleaned with the CFC composition. The ester composition functioned as a drop-in replacement for the CFC composition, with no harmful effects on the workers.

30 Since the flash point of the ester composition of Example 1 is 285°F, the potential for explosion or fire was not present as a result of the placement. However, a very significant

cost saving was achieved in the replacement because there were no significant evaporative losses of the ester composition and the time interval for change of the ester composition in the container was extended by at least three times. The latter  
5 advantage was realized because the ester composition continued to remove the chips and grinding residues from the parts even though the concentration of cutting oil in the ester composition was more than three times greater than in the CFC composition. A further cost saving was achieved by arranging  
10 for a blow-off station to catch the mist which was blown off the parts after they were dipped in the ester composition. The captured mist was collected and the liquid was reused in the cleaning process.

15 The next example is a comparative example showing the use of a prior art, commercially available cleaning composition comprising mineral spirits. This comparative example is followed by an example which also shows the use of the ester composition of Example 1.

Example C-2

20 This example also involved the cleaning of components parts of automotive fuel injectors. The parts were subjected to various machining operations and then placed in the solvent manifolds of HAPA Galvanotechnik washing machines where a stream of mineral spirits (solvent) impinged on the parts to remove the cutting oil, chips and grinding residues. The chips  
25 and grinding residues were filtered continuously from the solvent in the machine and the cutting oil concentration built up in the solvent as the solvent cycled through the machine. The solvent was replaced in the machines twice per week during  
30 operation. During the cleaning operation involving the use of mineral spirits, the operators complained of headaches, nausea and dermatitis.

Example 2

The mineral spirits used in the washing machines referred to in Example C-2 was replaced with the ester composition of Example 1. The fuel injectors were cleaned in the same machine cycles without complaints of headaches, nausea or dermatitis from the machine operators. The risk of explosion or fire was greatly reduced because the flash point of the ester composition is 285°F, as compared to 104°F for the mineral spirits. Furthermore, a drastic cost reduction was achieved because the solvent change-out frequency in the machines was extended from twice per week for the mineral spirits to once every two months for the ester composition. This was due to the ability of the ester composition to clean effectively even though loaded with about 10 wt.% cutting oil. Cost savings accrued also from other factors, such as reduced disposal costs, no evaporative losses and reduced machine down-time.

The next example is a comparative example which shows the use of a prior art, commercially available cleaning composition to clean transmission parts. The comparative example is followed by an example of the present invention showing also the effective use of a composition like that of Example 1. In these examples, residual cleaning composition was removed from the parts by blowing with a jet of air.

Example C-3

This example involved the cleaning of steel parts at the site of a warranty transmission rebuilder for the automotive industry. The cleaning composition comprised mineral spirits that was used in three different kinds of washing machines to clean transmission parts before reassembly. The consumption of the composition was at the rate of about 2,000 gallons per week. Along with headaches and nausea, 168 cases of dermatitis were reported during 12 months of use of the composition.

Example 3

The C-3 cleaning composition was replaced by the ester composition of Example 1. The transmission parts were cleaned with no reports of dermatitis, headaches or nausea. The 5 transmission parts were more effectively cleaned with the ester composition as determined by the Sediment Test, a standard test used in the automotive transmission industry to evaluate cleanliness. The Sediment Test values were reduced by more than 60% when the ester composition was used in place of the C-10 3 composition. The potential for fire or explosion was greatly reduced because the flash point of the ester composition is 285°F, as compared to 108°F for the C-3 composition. The consumption rate of the ester composition was 108 gallons per week whereas that of the C-3 composition was 2,000 gallons per 15 week. Even though the ester composition is more costly than the C-3 composition, a cost savings of more than 58% was achieved in the conversion to the ester composition because of its reduced rate of consumption. The reduced consumption of the ester composition came about because it continued to clean effectively even though loaded with about 15 wt.% of residual oil.

The next example is a comparative example which shows the use of a prior art, commercially available CFC composition comprising about 94 wt.% 1,1,2-trichlorotrifluoroethane and about 6 wt.% methanol to clean solder flux of the rosin type from electrical parts. The comparative example is followed by 25 an example which shows the use of a composition of the present invention to clean the aforementioned type of parts.

Example C-4

In this example, parts for small rotary electrical switches were made by soldering a small brass disk onto a small brass shaft. The solder flux, which congealed and hardened at 30

the solder joint, was removed from the resulting part using a cleaning composition comprising 1,1,2-trichlorotrifluoroethane, that is, a halogenated solvent of the type that is associated with ozone depletion. The parts were cleaned by immersing them 5 in the cleaning composition which was contained in a heated ultrasonic vat equipped with a vented hood exhaust system. The composition was removed by air drying.

Example 4

10       The composition of Example C-4 was replaced by a composition comprising approximately 90 wt.% isopropyl myristate and approximately 10 wt.% surfactant composition (Tergitol NP-4 (3.33 wt.%), Tergitol NP-7 (1.66 wt.%), Macol 2 (1.66 wt.%), Cyclogol SBDO (1.66 wt.%), and dipropylene glycol 15 (1.66 wt.%). The solder flux was removed from the parts every bit as well with this composition as with the composition of Example C-4. The residual ester-based composition was removed from the parts by rinsing with tap water. No complaints of any health or safety problems were reported in the conversion to 20 the use of the ester-based composition. Since the flash point of the ester-based composition is 285°F, the potential for fire or explosion was minimal. Very significant cost savings were achieved in the conversion from the composition of Example C-4 to the ester-based composition.

25       The next example is a comparative example which shows the use of a prior art, commercially available halogenated hydrocarbon solvent to clean brass parts. The comparative example is followed by an example which shows the use of a composition like that of Example 4 above to clean the same type 30 of brass parts.

Example C-5

In this example, brass parts were fabricated by turning in high production screw machines. The cutting oil, chips and turnings, were removed from the fabricated parts in a vapor degreaser using 1,1,1-trichloroethane. The composition was removed from the brass parts by air drying.

Example 5

The cleaning process of Example C-5 was changed to one which provided for placing the fabricated parts from the screw machines into wire mesh baskets and then dipping the baskets of parts repeatedly in a container of the cleaning composition like the composition of Example 4. The composition effectively removed the cutting oil, chips and turnings from the parts. The residual composition was removed from the parts by rinsing with water. The ester-based composition was very well received by the operators handling and using it. No complaints of any health or safety problems were reported. Cost savings in the conversion were significant.

The next three examples are illustrative of the practice of the present invention and show some additional applications in which monobasic esters can be used effectively.

Example 6

Methyl laurate (100%) was used to clean small metal parts of metal chips and cutting oils. The parts were dipped in the methyl laurate, blasted with compressed air to remove excess liquid and soils, and gauged with measuring instruments to determine part dimensions. The results were deemed to be fully satisfactory relative to the use of a conventional chlorofluorocarbon (CFC-113).

Example 7

In this example, soiled laboratory uniforms were dry cleaned in a cleaning composition comprising 88 wt.% isopropyl myristate, 12 wt.% of a surfactant mixture consisting of 5 Tergitol NP-4 (4 wt.%), Tergitol NP-7 (2% wt.%), Macol 2 (2 wt.%), cyclogol SBDO (2 wt.%), and dipropylene glycol (2 wt.%). Following the wash, the uniforms were rinsed in water. After drying, the uniforms were found to be satisfactorily clean.

Example 8

10 In this example, a cleaning composition comprising the methyl ester of capric acid was used to clean a printed wiring assembly which had been soldered using rosin flux. The cleaning performance was equivalent to that obtained with a terpene/surfactant mixture (BIOACT<sup>®</sup> EC-7<sup>TM</sup>, Petroferm Inc.).

15 Thus, monobasic esters are useful as cleaning agents. They are capable of dissolving a variety of common contaminants and offer the advantages indicated earlier relative to other cleaning agents.

Terpene Development/Use of Monobasic Ester Preferred

20 An important characteristic of the present invention is that it is capable of being used effectively in an industrial cleaning process, as distinguished from a cleaning process that is simply carried out on a laboratory scale or a pilot-plant scale. The invention can be practiced in either a batch or 25 continuous cleaning process.

Although it is expected that the invention will be used most widely in cleaning printed circuit boards, it has wider applicability in that it can be used to clean any type of substrate which is contaminated with soldering flux and/or 30 other types of residue that typically contaminate the surfaces

of printed circuit boards, for example, adhesive tape residue, waxes and fingerprints. Thus, printed wiring boards and other articles which require the use of soldering flux in the making of mechanical, electromechanical or electronic connections with metal can be cleaned in accordance with the present invention. In the fabrication of such articles, rosin soldering flux is used widely. However, there also can be used other types of soldering fluxes, including, for example, synthetic activated fluxes, organic acid-based fluxes and fluxes contained within soldering paste.

The terpene-based cleaning composition for use in the practice of the present invention comprises at least about 60 wt.% terpene, more preferably at least about 80 wt.% terpene, and most preferably about 85 to about 95 wt.% terpene. The composition can comprise about 100 wt.% terpene, but preferably is used in admixture with another constituent as described hereinbelow.

The terpene for use in the practice of the present invention is a terpene compound or a mixture of terpene compounds that are substantially water-insoluble and that are capable of dissolving the involved contaminants. The use of such terpene to clean printed circuit boards and related types of articles of contaminants is well known. For practice of the present invention, the use of para-menthadienes is preferred, with dipentenes being more preferred and limonene being most preferred. Excellent results have been achieved utilizing a mixture of terpene in which about 90 to about 95 wt.% of the mixture is limonene (the major portion being d-limonene), with the balance of the mixture being other terpene compounds.

A particularly preferred embodiment of the present invention comprises a terpene-based composition which includes also a monobasic ester, which is a material that is also substantially water insoluble, as mentioned hereinabove. Such

esters generally comprise the reaction product of alkyl alcohols and fatty acids, for example, C<sub>1</sub> to about C<sub>6</sub> alkanols and about C<sub>8</sub> to about C<sub>18</sub> fatty acids. Preferably C<sub>1</sub> to C<sub>4</sub> alcohols are used. Preferred monobasic esters are methyl laurate, isopropyl laurate and methyl caprate.

The monobasic ester can comprise up to about 40 wt. % of the terpene-based composition, but preferably comprises up to about 20 wt. %, most preferably about 5 to about 15 wt. % of the composition. Particularly good results having been achieved with the use of about 10 wt. % of the ester.

It should be understood that a material different from a monobasic ester (an optional, but preferred ingredient) can also be included in the terpene-based cleaning composition. Such material is desirably one which meets the COD standards set forth herein. As mentioned above, the terpene-based cleaning composition is substantially free of water-soluble material, that is, according to the procedure described above, the COD of the water phase of a two-phase mixture of equal amounts of the terpene-based cleaning composition and water is not greater than about 1000 ppm.

The contaminated surface should be contacted with the cleaning composition for a period of time sufficient to solubilize the contaminant. The time for effecting solubilization will depend on the temperature of the cleaning composition and the means by which it is applied. The temperature of the composition can be like that typically used for terpene-based compositions in applications of the present type, for example, room or ambient temperature (about 70°F) to about 150°F. Exemplary means for applying the cleaning composition are immersion, brushing, and spraying, for example, spraying in air or beneath the surface of the cleaning composition. It is believed that spraying the cleaning composition onto the substrate will be used most widely. For

batch operations, typical contact times are about 3 to about 5 minutes. For continuous operations, typical contact times are about 30 seconds to about 2 minutes.

- The solubilized contaminant is removed from the terpene-treated substrate by contact with water having a temperature of about 70°F to about 140°F. Good and satisfactory results have been obtained utilizing rinse water at room temperature. Water-contact times of the type generally used in the art can be used, for example, not longer than about 10 minutes.
- 10 Preferably, the water-contact time does not exceed about 5 minutes. Good results have been achieved with a contact time of about two minutes.

Conventional means can be used to apply the water. It is believed that, for most applications, spraying the air-borne substrate will be satisfactory and convenient. A spray pressure of about 30 to about 50 psi is exemplary. Other means for applying the water can be used also, for example, ultrasonic water agitation.

20 After the solubilized contaminants have been removed from the substrate, the substrate is dried. Typically, drying will be accelerated by the use of heat. For example, the substrate can be dried in an oven and/or it can be dried with forced air, for example, heated to 100°F or above.

25 With respect to the degree of substrate cleanliness that can be achieved by the practice of the present invention, levels of contamination that meet or are below the military specification MIL-P-28809A requirement of 14 micrograms NaCl equivalent/square inch can be achieved. The invention has been used effectively to provide substrates having a contamination level below about 10 micrograms NaCl equivalent/square inch.

It should be understood that such low levels of contamination can be achieved without the use of a surfactant in the water which is used to remove the solubilized contaminant from the substrate. Indeed, the COD of such water 5 should be such as to meet the COD requirements set forth herein. Deionized water is used preferably.

The effluent produced by the cleaning process comprises a liquid mixture of the terpene-based cleaning composition, contaminant dissolved therein, and water. The liquid mixture 10 may contain some solids, for example, minute pieces of solder and miscellaneous debris. The nature of the liquid mixture produced in accordance with the present invention is such that the components thereof can be separated conveniently by gravitational force. Thus, upon being fed to a settling tank, 15 solids precipitate from the mixture and the liquid components thereof stratify into a top layer comprising the water insoluble or immiscible liquids, that is, terpene and organic materials dissolved therein, including contaminants, and an aqueous bottom layer. The nature of the process is such that the aqueous layer is substantially free of water-soluble 20 organic material, that is, it has a COD of no greater than about 1000 ppm. Indeed, the process of the present invention can be utilized effectively to provide an aqueous phase in which the COD thereof is about 300 ppm or less, for example, 25 no greater than about 150 ppm. And, this can be accomplished without subjecting the water to any purifying steps or dilution.

In separating the different phases of the liquid mixture, the top organic layer can be simply drawn off. If desired, the aqueous phase can then be discharged directly into the sewage 30 system, while meeting the COD demands imposed by the involved governmental agency. If desired, the aqueous phase can be recycled for reuse in the cleaning process. This will generally involve the steps of filtration to separate solids

and ion exchange to remove undesirable inorganic materials, the latter step being preceded by carbon adsorption (if necessary) to remove organic materials that may adversely affect the ion exchange step. Other means may be used to purify the water.

- 5       Other means can be used to separate the liquid phases of the mixture, for example, centrifugation or by use of appropriate membranes.

10      The terpene compounds can be treated for recycling or they can be disposed of in an appropriate way, for example, incineration.

#### Examples of Terpene Development

This example is illustrative of the practice of the present invention.

##### Example A

- 15      A set of three identical printed circuit boards was fluxed with a rosin soldering flux (ALPHA 627 flux sold by Alpha Metals Inc. of Jersey City, New Jersey), wave soldered, and then cleaned of the flux residue that was adhered to each of the boards.
- 20      The cleaning composition contained terpene which were derived from orange peels and which comprised about 95 wt.% limonene and 5 wt.% of a mixture of other terpene compounds. The cleaning composition was substantially free of water-soluble organic materials. The boards were placed in the rack of an industrial dishwasher (semi-aqueous cleaning module) sold by Electronic Controls Design as Model 6307. The cleaning operation involved spraying the boards for about 5 minutes with the cleaning composition which was at room temperature (about 75°F).
- 25

The terpene-treated boards were then rinsed with water to remove the cleaning composition and flux dissolved therein. This was accomplished by subjecting the boards to a 1-minute, 20-second water wash in a continuous process carried out in an 5 Electrovert H-500 water rinse unit sold by Electrovert of Arlington, Texas. The spray pressure was about 40 pounds/square inch. The temperature of the water was about 120°F. The water collected from the rinsing operation has a COD well below 1000 ppm. Thereafter, each of the cleaned 10 boards was dried with hot air.

The cleanliness of each of the boards was evaluated by use of an Omega meter 600 SMD ionic testing device (sold by Alpha Metals Inc.). The average for the three boards was a contamination reading of only 9.3 micrograms NaCl 15 equivalent/square inch.

The next example is a comparative example which is illustrative of a prior art process.

Example No. C-A

The same procedure as described in Example No. 1 above was 20 followed except that cleaning was effected with a composition comprising 90 wt.% of the terpene cleaning composition described in Example 1 above and 10 wt.% of a mixture of emulsifying surfactants. The mixture comprised 50 wt.% of an ethoxylated sorbitan ester and 50 wt.% of an ethoxylated linear 25 alcohol. The average for the three boards that were cleaned was a reading of 9.4 micrograms NaCl equivalent/square inch - essentially the same as that of the boards of Example No. 1. However, the rinse water has a COD in excess of 5,000 ppm.

Claims

1. An industrial cleaning process in which a contaminant comprising soldering flux and/or adhesive tape residue is removed from a substrate contaminated therewith comprising:
  - 5 (A) contacting the substrate with a terpene-based cleaning composition which is substantially free of water-soluble organic material for a period of time sufficient to solubilize the contaminant;
  - 10 (B) removing the composition and solubilized contaminant from the substrate by contact with water having a temperature of about 70°F to about 140°F for a period of time of no longer than about 10 minutes to provide a substrate having a contamination rating of no greater than about 14 micrograms NaCl equivalent/square inch (MIL-P-28809A), and to
  - 15 thereby form a mixture comprising the composition, contaminant and water; and
- 20 (C) separating the water from said mixture, the separated water being substantially free of water-soluble organic material and having a chemical oxygen demand of no greater than about 1,000 ppm.
2. A process according to Claim 1 wherein the composition and solubilized contaminant are removed from the substrate by water spray.
- 25 3. A process according to Claim 1 wherein said contact with water is effected under conditions to provide a substrate having a cleanliness rating of below about 10 micrograms.
4. A process according to Claim 1 wherein the terpene-based composition comprises at least about 80 wt.% of terpene compound.

5. A process according to Claim 4 wherein the terpene-based composition comprises about 85 to about 95 wt.% of terpene compound.
- 5 6. A process according to Claim 5 wherein the composition comprises about 5 to about 15 wt.% of monobasic ester.
7. A process according to Claim 6 wherein the ester is methyl laurate, isopropyl laurate, or methyl caprate.
8. A process according to Claim 1 wherein the water is separated from the mixture by gravitational force.
- 10 9. A process according to Claim 8 wherein the separated water is discharged to a sewage system.
- 10 10. A process according to Claim 9 wherein the COD of the separated water is no greater than about 300 ppm.
- 15 11. A process according to Claim 10 wherein the COD of the separated water is no greater than about 150 ppm.
12. A process according to Claim 1 wherein the substrate is contaminated with soldering flux.
13. A process according to Claim 1 wherein the flux is rosin flux.
- 20 14. A process according to Claim 1 wherein the temperature of the water is about 110°F to about 130°F and the period of time is no longer than about 5 minutes.
15. A method for removing an organic residue from a surface contaminated therewith comprising:

- (A) contacting the surface with a cleaning composition comprising a major amount of a monobasic acid ester;
- 5 (B) maintaining the cleaning composition in contact with the surface for a period of time at least sufficient to solubilize the residue; and
- (C) removing cleaning composition and solubilized residue from the surface.

16. A method according to Claim 15 wherein said composition is used to clean a plurality of parts having their surfaces contaminated with said residue and wherein said residue is dissolved by said composition and wherein said dissolved residue builds up in concentration in said composition as the composition is used to clean additional parts and continuing to use said composition to clean additional parts when the concentration of the dissolved residue is about 20 to about 40 wt.%.

15

17. A method according to Claim 16 including continuing to use said composition to clean additional parts when the concentration of the dissolved residue is about 20 to about 30 wt.%.

20

18. A method according to Claim 15 wherein said monobasic ester has the formula



25 wherein R is an alkyl group having 1 to about 4 carbon atoms and R<sub>1</sub> is an alkyl group having about 8 to about 16 carbon atoms.

19. A method according to Claim 15 wherein the monobasic ester is methyl laurate, isopropyl laurate, methyl caprate, or isopropyl myristate

30 20. An organic-based composition comprising a major amount of a monobasic ester and a minor amount of another cleaning agent selected from the group consisting of terpenes,

esters, dibasic esters, petroleum hydrocarbons, ketones, ethers and alcohols.

21. A composition according to Claim 20 comprising about 55 to about 90 wt.% of th monobasic ester and about 10 to 5 about 45 wt.% of said other cleaning agent.

22. A composition according to Claim 20 including also a surfactant.

23. An organic-based cleaning composition comprising a major amount of a monobasic ester and about 2 to about 20 wt.% 10 surfactant.

24. A composition according to Claim 23 wherein the surfactant comprises about 8 to about 12 wt.% of the composition.

25. A composition according to Claim 23 wherein the ester 15 is either methyl laurate, isopropyl laurate, methyl caprate or isopropyl myristate and said surfactant is an ethoxylated alkylphenol or a detergent alcohol.

# INTERNATIONAL SEARCH REPORT

International Application No. PCT/US91/08937

## I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) \*

According to International Patent Classification (IPC) or to both National Classification and IPC  
 IPC(5): B08B 7/00 C11D 7/50  
 U.S. 134/40

## II. FIELDS SEARCHED

Minimum Documentation Searched ?

Classification System	Classification Symbols
U. S.	134/40, 134/42, 252/162, 252/170, 252/Dig. 14

Documentation Searched other than Minimum Documentation  
to the Extent that such Documents are Included in the Fields Searched \*

APS Search: TERPENE, MONOBASIC ACID ESTER

## III. DOCUMENTS CONSIDERED TO BE RELEVANT \*

Category *	Citation of Document, ** with Indication, where appropriate, of the relevant passages †‡	Relevant to Claim No. †‡
P, Y	US, A, 4,983,224 (MOMBRUN ET. AL.) 08 JANUARY 1991 See claims and Col. 5.	1-5 & 8-14
Y	US, A, 4,867,800 (DISHART ET. AL.) 19 SEPTEMBER 1989 See claims.	1-5 & 8-14
A	US, A, 4,934,391 (FUTCH ET. AL.) 19 JUNE 1990	
A	US, A, 4,927,556 (POKORNY) 22 MAY 1990	
A	US, A, 4,740,247 (HAYES ET. AL.) 26 APRIL 1988	

\* Special categories of cited documents: †‡

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

## IV. CERTIFICATION

Date of the Actual Completion of the International Search

14 FEBRUARY 1992

Date of Mailing of this International Search Report

11 MAR 1992

International Searching Authority

ISA/US

Signature of Authorized Officer

Saeed T. CHAUDHRY